

L-38608-65

ACCESSION NR: AP5005310

S

possible to measure the birefringence produced in the crystal. An electron optical converter was used to change the infrared radiation into visible light. The results yielded the dispersion of the photoelastic constants of gallium arsenide and of silicon for the near infrared region of the spectrum. The photoelastic constants were observed to be strongly dependent on the wavelength, with especially strong dispersion at the absorption edge. This wavelength dependence cannot be attributed to the dispersion of the refractive index but is a feature of semiconductors. "The authors thank M. V. Klassen-Neklyudova and V. L. Indenbom for support and a discussion of the results, and L. N. Mikheylov and A. I. Bober for supplying the material necessary for the research." Orig. art. has: 1 figure and 4 formulas.

ASSOCIATION: Institut fiziki tverdogo tela AN SSSR, Moscow (Institute of Solid State Physics, AN SSSR)

SUBMITTED: 16Jun64

ENCL: 00

SUB CODE: SS, OP

NR-REF BOV: 002

OTHER: 006

Card 2/2

L 08398-67 EWT(n)/EWP(t)/ETI IJP(c) JD/JG SOURCE CODE: UR/0069/66/028/005/0692/0695  
ACC NM AP6032178

AUTHOR: Martynenko, G. P. (Moscow); [Malyshev, N. I.] (Deceased; Moscow)

ORG: none

TITLE: Adsorption of copper from aqueous solutions on the surface of gallium arsenide

SOURCE: Kolloidnyy zhurnal, v. 28, no. 5, 1966, 692-695

TOPIC TAGS: gallium arsenide, adsorption, copper

ABSTRACT: The adsorption of copper ions on polished gallium arsenide from aqueous HCl, KOH, H<sub>2</sub>SO<sub>4</sub>, and H<sub>2</sub>O solutions was studied by using the Cu<sup>64</sup> radioisotope tracer. The absorption isotherms were determined, and it was found that the adsorption increases with decreasing pH of the solution. Maximum adsorption was observed in the case of KOH. At a Cu<sup>64</sup> concentration of 10<sup>-7</sup> g/ml in the alkaline solution, the adsorption amounted to 5 x 10<sup>-8</sup> g/cm<sup>2</sup>. In all cases (H<sub>2</sub>O, acidic and alkaline solutions) the adsorption was irreversible. The distribution of copper over the surface of gallium arsenide was determined by radiography and showed that the adsorption is most pronounced in pits, cracks, scratches, fused metal contacts, and p-n junctions. In order to minimize contamination with copper, acid rather than alkaline etchants are recommended for treatment of gallium arsenide surfaces. Orig. art. has 3 figures.

SUB CODE: 07,20/ SUBM DATE: 24Mar65/ OTH REF: 006

Card 1/2 afs

UDC: 541.183.24

MARTYNEKO, I.A.

Using the ShM-2 cutter-loader for rapid drifting in the "Velikomos-tovskaiia" Mine No.3. Ugol' 35 no.8:11-12 Ag '60. (MIRA 13:9)

1. Glavnnyy inzhener tresta Novovolynskugol'.  
(Lvov-Volyn' Basin--Coal mining machinery)

MIKHAYLOV, V.G., doktor tekhn.nauk; KHAIIVIL, M.G., kand.tekhn.nauk;  
KARYUK, G.G., kand.tekhn.nauk; KAZHENTSEV, Yu.T., aspirant;  
GARASHCHENKO, P.A., aspirant; MALYAKOV, G.P., aspirant;  
KOGAN, K.b., inzh.; SUKACH, V.D., inzh.; TKACHENKO, V.A., inzh.;  
LINEENKO, Yu.F., inzh.; MOZNAIM, G.I., inzh.; MARTYNENKO, I.A., inzh.

Cutting tool for the cutter loader. Ugol' Ukr. 6  
(MIRA 15:11)  
no.8:37-39 Ag '6x.  
(Coal mining machinery)

MARTYNNENKO, I.A., inzh.; MILEYAYEV, I.S., inzh.; TUGAYEV, T.S., inzh.;  
KOTLYARSKIY, I.A., inzh.; MOREV, A.B., inzh.; MIDRYAK, V.A.,  
inzh.; SUDOPLATOV, A.P., prof.; IVANOV, K.I., kand. tekhn. nauk;  
IGNAT'YEV, A.P., kand. tekhn. nauk; KOLYSKIN, G.M., kand. tekhn.  
nauk; YEREMENKO, Ye.I., inzh.

Industrial testing of the auger drilling of coal with double  
spindle auger drilling machines. Ugol' 40 no.1:32-37 Ja '65.  
(MIRA 18:4)

1. Kombinat 'Dkrzapadugol' (for Martynenko, Miliyayev, Tugayev).
2. Gerlovskiy mashinostroitel'nyy zavod im. S.M.Kir'ya (for Kotlyarskiy, Morev, Midryak). 3. Institut gornogo dela im. A.A.Skochinskogo (for Sudoplatov, Ivanov, Ignat'yev, Kolyskin, Mel'nikov, Yeremenko).

MARTYNNENKO, I.A.; PROTOPOPOV, S.F.

New developments in working twin long walls. Ugol' 40 no.9:18-20  
(MIRA 18:10)  
S '65.

1. Glavnyy inzh. tresta Novovolynskugol' (for Martynenko).
2. Zamestitel' glavnogo inzhenera shakhty No.5 "Novovolynskava"  
(for Protopopov).

44-839  
S/560/62/000/014/011/011  
A001/A101

AUTHORS: Krylov, G. N., Martynenko, I. A., Pogrebnyak, Ye. B., Sergeyeva, M.K.

TITLE: An autonomous optical method of determining orientation of an Earth's satellite in space

SOURCE: Akademiya nauk SSSR. Iskusstvennye sputniki Zemli. no. 14, 1962,  
145 - 153

TEXT: The purpose of this investigation is determination of orientation of an Earth's artificial satellite in space from the readings of solar radiation pickups mounted on the satellite. In the first part of the article the problem of determining the vector, denoting direction from the satellite to the Sun, is solved in terms of three direction cosines. The second part deals with the problem of determining the matrix of transformation from the coordinate system "Earth" to the system of "Sputnik". As a result, the matrix was obtained, each of whose elements contained the unknown angle of turn of one system relative to the other. This angle can be determined from the reading of at least one scanning pickup whose optical system fixes the line of horizon, provided that the altitude of the

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An autonomous optical method of...

S/560/62/000/014/011/011  
A001/A101

satellite above the Earth's surface is known. If the readings of several scanning pickups are known, which determine the line of horizon, then it is possible to calculate from them both the angle of turn and the altitude of the satellite. In the fourth part of the article the authors analyze the problem of forecasting the orientation of the satellite in space. For this purpose they derive the formula for the angular velocity of the satellite using three Euler dynamical equations of rotational motion and three kinematic equations. Then, knowing the angular velocity and time intervals expired from the beginning of motion, one can calculate for the future the orientation of the satellite in space. In conclusion the authors discuss the problem on the number of pickups necessary for determination of orientation; the number is eight, located at the vertices of a cube, of which four will be illuminated by the Sun. There is 1 figure.

SUBMITTED: February 26, 1962

Card 2/2

MAR' YNENKO, I.A., na valokreuil'shchik

Produce motion pictures on safety measures for Moscow Basin. Bezou.  
truda v prom. 2 no.5:35 My '58. (MIRA 11:4)

1. Shakhhta No.37 tresta Stalinogorskugol', predsedatel' uchastkovogo  
komiteta profsoyuza rabochikh ugol'noy oromyshlennosti.  
(Moscow Basin--Motion pictures in industry)

MARTYNENKO, I. I.

"Operation of Three-Phase Asynchronous Motors of Agricultural Electrical Equipment With Nonsymmetry of Voltages." Cand Tech Sci, Moscow Inst of Electrification and Mechanization of Agriculture imeni V. M. Molotov, Min Higher Education USSR, Moscow, 1954. (KL, No 1, Jan 55)

Survey of Scientific and Technical Dissertations Defended at USSR Higher Educational Institutions (12)  
SO: Sum. No. 556, 24 Jun 55

MARTYNNENKO, I.I., kand.tekhn.nauk

Allowed load for electric motors operating under voltage deviation.  
Mekh. i elek.sots.sel'khoz. no.4:40-42 '57. (MIRA 12:4)

1.Ukrainskiy nauchno-issledovatel'skiy institut mekhanizatsii i  
elektrifikatsii sel'skogo khozyaystva.  
(Electric motors, Induction)

GLUSHCHENKO, V.P. [Hlushchenko, V.P.], kand.tekhn.nauk; MARTYRENKO, I.I.,  
kand.tekhn.nauk

Magneto as the source supplying electric fences. Mekh. sil'. hosp.  
[9] no.5:27-28 My '58. (MIRA 11:6)  
(Electric fences)

AUTHOR: Martynenko I.I., Candidate of Technical Sciences

TITLE: A Graphical Method of Determining the Working Characteristics of an Induction Motor with Asymmetrical Voltage (Opravleniye rabochikh kharakteristik asinkhronnogo dvigatelya pri nesimmetrii napryazheniy graficheskim sposobom)

PERIODICAL: Vestnik Elektropromyshlennosti, 1958, Vol 29, No 5, pp 29 - 31 (USSR).

ABSTRACT: Available analytical methods of determining the operating characteristics of induction motors with asymmetrical voltage are cumbersome and laborious. This article describes a graphical method of solving the problem, using the ordinary circle diagram with some additional constructions. The method is based on the theory of symmetrical components. The negative phase-sequence impedance is practically constant when the slip is small. Therefore, if the negative phase-sequence voltage is constant, the corresponding current is constant in value and direction.

The proposed circle diagram shown in Figure 1 is based on a simplified equivalent circuit of an induction motor with asymmetrical voltage. The usual circle diagram is constructed for positive phase-sequence currents corresponding to the positive phase-sequence voltages. The vectors of the actual

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10-58-52-70

A Graphical Method of Determining the Working Characteristics of an  
Induction Motor with Asymmetrical Voltage

phase voltages are obtained by geometrical summation of the vectors of negative and positive phase-sequence voltage. The vectors that join the start of the positive phase-sequence vectors to the ends of the negative phase-sequence vectors are the actual phase-voltage vectors. The relative positions of the vectors are not, however, correct and a method of determining the correct position is given. Next the method of determining the phase currents is described. The circle diagram is then constructed as shown in Figure 2 and the procedure for deriving currents and phase angles is explained.

In order to determine the validity and accuracy of the graphical methods, experiments were made on a series A-motor. Using a known asymmetric voltage supply, measurements were made of current and power and phase angles of voltage and current were calculated. A graphical construction was made for the same conditions. Comparison of the experimental and calculated results given in Figures 3, 4 and 5 show that the difference is comparatively small, being of the order of 5 - 10%.

Card 2/3

1.0-58-113/27

A Graphical Method of Determining the Working Characteristics of an Induction Motor with Asymmetrical Voltage

There are 5 figures and 6 references, 4 of which are Soviet and 2 English.

ASSOCIATION: Ukrainskiy nauchno-issledovatel'skiy institut mechanizatsii i elektrifikatsii sel'skogo khozyaystva  
(Ukrainian Scientific Research Institute for the Mechanisation and Electrification of Agriculture)

SUBMITTED: June 10, 1959

Card 3/3

MARTYNNENKO, Ivan Ivanovich, kand.tekhn.nauk; BOHDARENKO, S.P., kand.  
tekhn.nauk, glavnyy red.

[New developments in rural electrification] Nove v elektrifika-  
tsii sil'skogo naseleniya. Kyiv, 1959. 30 p. (Tovar-  
istyvo dlia poshyreniya politychnykh i naukovykh znanii Ukrainskoj  
ins'koi RSR. Ser.6, no.13).  
(Rural electrification)

BUDZKO, I.A., akademik, otv.red.; BONDARENKO, S.P., kand.tekhn.nauk,  
zamestitel' otv.red.; MARTYNEJKO, I.I.; KARPOV, I.V., red.;  
OLEYNIK, V.S., red.; KOSOVSKIY, V.A., red.; KVITKA, S.P..  
khudozhestvenno-tekhn.red.

[Problems connected with electric power supply to agriculture;  
collection of articles on materials of the scientific session  
of the section of the electrification of agriculture] Voprosy  
elektrosnabzheniya sel'skogo khoziaistva; sbornik statei po  
materialam nauchnoi sessii sektsii elektrifikatsii sel'skogo  
khoziaistva. Kiev, Izd-vo Ukr.akad.sel'khoz.nauk, 1959. 149 p.  
(MIRA 13:2)

1. Kiyev. Ukrains'ka akademija sil's'kohospodars'kykh nauk.
2. Vsesoyuznaya akademija sel'skokhozyaystvennykh nauk imeni V.I.  
Lenina, direktor Vsesoyuznogo nauchno-issledovatel'skogo instituta  
elektrifikatsii sel'skogo khozyaystva (VIESKh) (g.Moskva) (for  
Budzko).

(Electricity in agriculture)

MARTYNEKO, I.I., kand.tekhn.nauk

How to organize the charging of accumulators on the collective  
farm. Mekh.sil'.hosp. 8 no.9:32-3 of cover S '59.  
(MIRA 13:1)

(Storage batteries)

MARTYRENKO, Ivan Ivanovich, kand.tekhn.nauk; BONDARENKO, S.P.,  
kand.tekhn.nauk, otd.red.; GUHENKO, V.A. [Hurenko, V.A.].  
red.

[Operating electric equipment on collective and state farms]  
Ekspluatatsiia elektricheskikh i radhospakh.  
Kyiv, 1960. 31 p. (Tovarystvo dlia poshyrennia politychnykh i  
naukovykh znan' Ukrains'koi RSR. Ser.6, no.22). (MIRA 14:2)

(Electricity in agriculture)

MARTINENKO, I.I. [Martynenko, I.I.], kand.tekhn.nauk; TERPILO, M.M.,  
inzh.-elektrik  
Electric water heating in AGK-12 automatic waterers. Mekh.sil'.  
hosp. 11 no.1:29-30 Ja '60. (MIRA 13:4)  
(Water heaters)

MARTYNIENKO, I. I., kand.tekhn.nauk

Calculations for stator windings of electric motors. Mekh. sil'.  
hosp. 11 no. 7:29-30 J1 '60.  
(MIRA 13:10)  
(Electric motors)

MARTYNNENKO, I.I., kand.tekhn.nauk

Electric power plant can work without a reducer. Mekh. sil'. hosp.  
12 no. 2:31 F '61. (MIRA 14:4)  
(Electric power plants)

MARTYNNENKO, I.I., kand.tekhn.nauk; DOROSH, I.I., kand.ekonomicheskikh nauk

Electric power supply for collective and state farms in the  
Ukrainian S. S. R. Mekh. i elek. sots. sel'khoz. 19 no.6:45-48  
'61. (MIRA 14:12)

1. Ukrainskiy nauchno-issledovatel'skiy institut mekhanizatsii  
i elektrifikatsii sel'skogo khozyaystva (for Martynenko).
2. Ukrainskiy nauchno-issledovatel'skiy institut ekonomiki i  
organizatsii sel'skogo khozyaystva (for Dorosh).  
(Ukraine--Electricity in agriculture)

MARTYNENKO, Ivan Ivanovich; DOROSH, I.Y.; KISTEN', G.Ya.  
[Kysten', H.IA.]; KOLOMIYETS', I.F.[Kolomiet's', I.F.];  
LEVITSKAYA, G.P.[Levits'ka, H.P.], red.; GULENKO, O.I.  
[Hulenko, O.I.], tekhn. red.

[Use of electric power on the "Shliakh do komunizmu"  
Collective Farm] Vykorystannia elektroenergii v kolhospiv  
"Shliakh do komunizmu." Kyiv, Derzhsil'hospvydav UkrSSR,  
1962. 58 p.

(Electricity in agriculture)

DOROSH, I.Y. [Dorosh, I.I.], kand.ekonom.nauk MARTYENKO, I.I. [Martynenko, I.I.], kand.tekhn.nauk

Effectiveness of the use of electric power in agriculture. Tekh. syl'.  
hosp 13 n. 7:2' -27 Jl 1962.  
(MIKA 17:3)

MARTINENKO, I.I. [Martynenko, I.I.], kand.tekhn.nauk

How to operate a three-phase electric motor by a single-phase circuit. Mekh. sil'. hosp. 13 no.9:29-30 S '62. (MIRA 17:2)

MARTINENKO, I.I. [Martynenko, I.I.], kand.tekhn.nauk

How to adjust alternating current machines for electric welding.  
Mekh. sil'. hosp. 13 no.4:29-30 Ap '62. (MIRA 17; 3)

MARTYNENKO, I.I., kand. tekhn. nauk; TERPILo, M.M. [Terpylo, M.M.],  
starshiy nauchnyy sotrudnik

Automatic irradiation unit. Mekh. sil'. hosp. 14 no. 9:28  
S '63. (MIRA 17:1)

1. Ukrainskiy nauchno-issledovatel'skiy institut mekhanizatsii  
i elektrifikatsii sel'skogo khozyaystva (for Terpilo).

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CIA-RDP86-00513R001032610018-9"

DRENNOVА, K. A., prof.; GRISHIN, S. I., prof.; MARTYNEНKO, L. I.;  
DADAMUKHAMEDOV, A. N.; IBRAGIMOV, R. I.; AMILOVA, A. A.; FEL'DMAN, F. Ya.;  
MESHKOVA, N. P.; SHENKER, D. I.

Condition of the ears nose and throat in children of preschool age  
in Tashkent. Vest. otorin. no. 2: 60-62 '61. (MIRA 14:12)

1. Iz Otorinolaringologicheskoy kafedry (zav. - prof. K. A. Drennova)  
Tashkentskogo instituta usovershenstvovaniya vrachey.

(TASHKENT--OTOLARYNGOLOGY)

MARTYNNENKO, I.M.

Repair of worn-out fire grates. Energetik 2 no.3:10 Mr '54.  
(MLRA 7:5)  
(Furnaces--Grates)

MARTYNEJKO, I.M.

Improvement of the BTSR-1 chain grates. Sakh.prom. 28 no.4:32-33  
'54. (MLRA 7:7)

1. Oktyabr'skiy sakharnyy zavod.  
(Sugar industry--Equipment and supplies)

BARTOSHEVICH, Ye.N.; TSUKER, M.B.; LESHCHINSKAYA, Ye.V.; SOKOLOVA, I.S.;  
MARTYNNENKO, I.N.; ANDREYEVA, L.S.; ASHMARINA, Ye.Ye.

Poliomyelitislike paralytic diseases in children inoculated  
with live Sabin vaccine. Vest. AMN SSSR 18 no.6:16-21 '63.  
(MIRA 17: 1)

NAUMENKO, Yu.I.; MARTYNNENKO, I.N.

Bioelectric activity of muscles in tetanus patients. Zhur. nevr. i psikh. 64 no.9:1310-1315 '64. (MIRA 17:12)

1. Institut nefrologii (direktor - prof. N.V. Konovalov)  
AMN SSSR i Institut poliomiyelita i virusnykh entsefalitov  
(direktor - prof. M.P. Chumakov), Moskva.

MARTYNNENKO, I.N.

Date of birth: 01.01.1938. Present place of residence: 127055, Moscow, 10th paralellia  
district of Krymskaya, 10th floor, 10th apartment, phone: 5-00-00.  
(MIRA 17/12)

1. Post: post office worker, 1st class, chief supervisor (direktor).  
Residence: 127055, Moscow, 10th floor, 10th apartment, M.P. Chumakov, AMN SSSR, Moscow.

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— 1 —

and can, if necessary, be used in the same way.

1. *Chlorophytum comosum* (L.) Willd. var. *variegatum* (L.) Kuntze

**APPROVED FOR RELEASE: 06/14/2000**

CIA-RDP86-00513R001032610018-9"

1. MARTYNENKO, K. D.
2. USSR (600)
4. Evaporation
7. Thermal exchange in a plan of multistage evaporation. Trudy Len. inst. pishch. orom. 1, 1949.

9. Monthly List of Russian Accessions, Library of Congress, April 1953, Uncl.

MARTYARENKO, Konstantin Dmitriyevich; SLAVYANSKIY, Aleksey konstantinovich,  
retsenzent; MIKHAYLOV, M. I., redaktor; NIKOLAYEVA, I. I., redaktor  
izdatel'stva; KORASIK, N.P., tekhnicheskiy redaktor

[Technical equipment of hydrolysis and sulfite and alcohol plants]  
Tekhnologicheskoe oborudovanie gidroliznykh i sul'fitno-spirtovykh  
zavodov. Moskva, Goslesbumizdat, 1956. 251 p. (MLRA 9:10)  
(Chemistry, Technical--Equipment and supply)  
(Alcohol) (Sulfite liquor)

KAPLAN, M.N.; MARTYNEJKO, K.D.; MOLOCHNY, R.M.

Standard plan of a hydrolysis furfural plant. Gidroliz.i lesokhim.  
prom. 12 no. 3:25-29 '59. (MIRA 12:6)

1. Giprogidroliz.  
(Hydrolysis) (Furaldehyde)

MARTYNNENKO, K.D.; KAPLAN, M.N.

Production of furrurole in Italy and France. Gidroliz.i  
lesokhim.prom. 12 no.6:26-29 '59. (MIRA 13:2)

1. Giprogidroliz.  
(Italy--Furaldehyde) (France--Furaldehyde)

MARTYNNENKO, Konstantin Dmitriyevich; KEYL', I.A., retsenzent;  
PAVLOV, B.T., retsenzent; MEL'NIKOV, N.P., red.;  
SARMATSKAYA, G.I., red. izd-va; VDCVINA, V.N., tekhn. red.

[Processes, apparatus, and equipment for wood hydrolysis and  
the manufacture of wood chemicals] Protsessy, apparaty i obo-  
rudovanie gidroliznogo i lesokhimicheskogo proizvodstv. Moskva,  
Goslesbumizdat, 1961. 444 p.  
(Wood--Chemistry)

(MIRA 15:5)

AGRONOMOV, S., inzh.-mekhanik, starshiy tekhnolog po svarke.; MARTYNNENKO, L.,  
tekhnolog po svarke.

Automatic building up of shafts using the spiral technique. Mor.  
fot 18 no.9:17-19 S '58. (MIRA 11:10)  
(Electric welding) (Shafting)

MARTYNEENKO, L., inzhener.

Results of innovators' work. Mias.ind.SSSE 28 no.4:57-58 '57.  
(MIRA 10:7)

1. Aktyubinskiy myasokombinat.  
(Meat industry--Equipment and supplies)  
(Lubrication and lubricants) (Compressors)

MARTYNENKO, L., inzh.

Quick freezers with direct ammonia evaporation. Mias.Ind.SSSR  
30 no.6:45-46 '59. (MIRA 13:4)

1. Aktyubinskiy myasokombinat.  
(Aktyubinsk--Cold storage)  
(Refrigeration and refrigerating machinery)  
(Meat, Frozen)

MARTYNIENKO, L., inzh.

Self-cleaning screen grinder. Miss.ind.SSSR 31 no.1:52 '60.  
(MIRA 13:5)

1. Aktyubinskiy myasokombinat.  
(Aktyubinsk--Sewage--Purification)

VOLKOV, B.V.; MARTYNNENKO, L.A.; KOCHUROVA, G.A.

Determination of nitrocyclohexane in sewage. Gig.i san. 26 no.1:  
62-63 Ja '61. (MIRA 14:6)

1. Iz Lisichanskogo filiala Nauchno-issledovatel'skogo i proyektnogo  
instituta azotnoy promyshlennosti i produktov organicheskogo sinteza.  
(CYCLOHEXANE) (SEWAGE)

VOLKOV, B.V.; MARTYNENKO, L.A.; KOCHUROVA, G.A.

Use of methanol for the regeneration of spent activated  
carbon in the purification of industrial sewage from organic  
compounds. Gig. i.san. 26 no.9:82-84 S '61. (MIR 15:3)  
(CARBON, ACTIVATED)  
(METHANOL)

MARTYNENKO, L.F.

Determining the weight of a function of magnitudes balanced by  
the knot method. Trudy Inst.fiz. i mat. AN BSSR no.2:267-271  
'57. (MIRA 12:1)  
(Nets (Geodesy)) (Weight functions)

MARTYNEKO, L.F.

Lateral refraction and its effect of the precision of the  
measurement of horizontal angles. Trudy Inst.fiz.i mat.ÄN  
BSSR no.3:244-249 '59. (MIRA 13:4)  
(Surveying)

SOKUR, Ivan Tarasovich; MARTYNEKO, L.I., red.; RAKHLINA, N.P.,  
tekhn. red.

[Injurious rodents and their control] Shkidlyvi hryzuny i  
borot'ba z nymy. Kyiv, Vyd-vo AN URSS, 1963. 93 p.  
(MIRA 17:3)

VSEKHSVYATSKIY, Sergey Konstantinovich; MARTYNNENKO, L.I., red.;  
REKES, M.A., tekhn. red.

[Sun and interplanetary space] Solntse i mezhplanetnoe  
prostranstvo. Kiev, Izd-vo AN USSR, 1963. 83 p.  
(MIRA 17:2)

MARTY'ENKO, L.I.; ZINTSOVA, Ye.S.; MAKAROV, V.N., KIZNETSOVA, M.N.;  
KONDRAT'YEVA, D.N.; SOVA, N.G.; TARANETS, V.I., TOMAREV, S.S.

Stratigraphy of the iron ore complex in the Yarlovets deposit.  
Sbor.nauch.trud.KGRI no. 71:24-2' '63. (MIRA 17:7)

LUK, Aleksandr Naumovich; MARTYNNENKO, L.I., red.

[Memory, cybernetics, and thinking] Pamiat', kibernetika, myslennia. Kyiv, Naukov dumka, 1964. 82 p.  
(MIRA 18:1)

LUK, Aleksandr Naumovich; MARTYNOV, V.I., red.

[Memory, cybernetics, and logical functions, memory cycles,  
mysteries. Tytuł, naukowy, 1974. — 12 p.]

(AIA 1974)

MARYNIAK, L. I.

"Certain Problems Connected With the Chromatographic Separation of Rare Earth Elements."

Rare earth elements (extraction, Analysis, Use), Presented by the Institute of Geochemistry and Analytical Chemistry Prof. J. I. MARYNIAK, Lublin, Poland.

(Chemical Faculty of the Masaryk State University, Dr. M. I. LINDNER).

5(2) Page 1 book information 800/1/27

Absentia mark 8001. Institut geokhimi i mineralogicheskoi khimii

Bol'shometal'nye elementy i polucheniye, smolu, primenenie (Rare Earth Elements, Extraction, Products, Analysis and Application) Moscow, Izd-vo Akad. Nauk, 1958. 331 p. 2,200 copies printed.

Ed. D. I. Rybachikov, Professor; Editorial Board: I. P. Al'mashin, Corresponding Member, USSR Academy of Sciences, N. N. Zaslavskiy, Doctor of Chemical Sciences, A. V. Egorov, Candidate of Technical Sciences, V. I. Ermakov, Doctor of Chemical Sciences, M. M. Sosulin, Candidate of Chemical Sciences, and Yu. G. Okhrabrova, Candidate of Chemical Sciences Eds. of Publishing House: D. S. Trifonov and T. G. Levit'ev. Ed.: S. G. Borovitsch.

**PURPOSE:** This book is intended for scientists, chemists, teachers and students of higher educational institutions, chemical and industrial enterprises and other persons concerned with the extraction, preparation, use or study of rare earth elements.

**CONTENTS:** This collection contains reports presented at the June 1956 Conference on Rare Earth Elements at the Institute of Geochemistry and Analytical Chemistry (Academy of Sciences USSR). The article deals with methods of separating rare earth elements, methods of processing rare earth ores, ion exchange chromatography, chemical analysis and some industrial applications of rare earths. Aside from contributing authors, the editors mention the following Soviet scientists who are studying rare earth elements: Parfenov, Mal'nikov, Churkin, Mol'ter, Matrosov, Chernyayev, Dan'yan, Mal'nikov, Zinov'ev and especially, I. A. Grigor'yev, who first obtained the molecular compounds of the rare earth elements in the pure state, separated many complex substances and won at the end of each article.

TABLE OF CONTENTS:

- Andreev, L.P., T.V. Libichenko, I.V. Andreev'eva, and O. I. Bratkovskaya (State Rare Metals Scientific Research Institute and Moscow Agricultural Academy Inst. E. A. Shchukarev), Trillium & in the Ion Exchange Separation of Less Common Rare Earth Elements 100
- Andreev, L.P., and A.S. Kostyug (Moscow Agricultural Academy Inst. E. A. Shchukarev and State Rare Metals Scientific Research Institute).
- Characteristics of Trillium & in the Ion Exchange Separation of Elements in the Cerium Subgroup 108
- Borzenko, L. I. (Moskovskiy gosudarstvennyy universitet imeni M.V. Lomonosova, Khimicheskii fakultet). Some Problems of the Chromatographic Separation of Rare Earth Elements 112
- Bordov, Yu. D., V. P. El'makov, and V. S. Golosova (Botanicheskii gordonetskiy universitet, Institut I. G. Chernysheva), Separative methods in the synthesis of lanthanide and actinide thorovanadates (Faculty of Chemistry, Institute of Chemistry and Physics Institute for the Mechanization of Agriculture in Odintsovo)
- Experiments on the separation of cerium isotopes in Ostankino 116

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(S)ensitive (Unclassified)

APPROVED FOR RELEASE: 06/14/2000

CIA-RDP86-00513R001032610018-9"

## AUTHORS:

Mitrofanova, N. L., Martynenko, L. I., S V. "D-1-11-11, 1"  
Yeremin, G. K.

## TITLE:

On Some Properties of the Complex Acids Produced From Rare  
Earths With Ethylene Diamine-Tetraacetic Acid (In ne-  
torykh svoystvakh kompleksnykh kislot, obrazovannykh  
reikozemel'nymi elementami i etilenidiamintetraakyskoy  
kisloty)

## PERIODICAL:

Zhurnal neorganicheskoy khimii, 1964, Vol 3, Nr 11, p. 2367-2368

## ABSTRACT:

The complex compounds of the rare earths (La, and Nd, Sm)  
diamino-tetraacetic acid ( $H_4V$ ) were investigated. The  
composition and the solubility of these complex acids  
were determined. Equivalent quantities of aqueous  
suspensions of  $H_4V$  were transferred with aqueous suspensions  
of the oxides of rare earths at room temperature. A pre-  
cipitate of compounds of the following composition is  
produced:  $H[LaV]$ . 6  $H_2O$ ,  $H[NdV]$ . 6  $H_2O$ ,  $H[SmV]$ . 6  $H_2O$ .  
At low temperature the cerium earths form complex compounds  
with crystal water of integral molar number. Anhydrous  
modifications of the complex acids with low solubility  
are produced from the boiling solutions. The formation  
Carri 1, 2

On Some Properties of the Complex Acids Prepared From Rare Earth Salts and Earths With Ethylene Diamine-Tetraacetic Acid

of the complexes is as well possible by means of the acidification of the complex salts of the type  $\text{M}^{\text{3+}}[\text{LnY}]_x \cdot \text{La}[\text{LnY}]_x$ . The yttrium oxide form as well  $\text{H}_3\text{Y}$  complex

complexes with different water content. Analysis is made which should not be produced in the case of yttrium oxides. The solubility of the complex acids of La, Pr, Nd and Sm with  $\text{H}_4\text{Y}$  at  $25^{\circ}\text{C}$  is given in table 2. On the strength

of the solubility difference between yttrium oxides and cerium earths a fractional separation via the adipic acid acids is suggested. There are 4 tables and 1 reference.

3 of which is Soviet.

ASSOCIATION. M. Skovskiy gosudarstvennyy universitet im. M.V. Lomonosova  
Kaletra neorganicheskoy khimii (Moscow State University  
imeni M.V. Lomonosova, Chair of Inorganic Chemistry,

SUBMITTER. September 1, 1977  
Car: 2, 2

AUTHOR: Martynenko, I. I. SOV/156-18-4-26/49

TITLE: The Determination of Ethylene Diamine Tetra Acetic Acid in the Presence of the Rare Earth Elements (Oprеделение этилендиаминететракисусульфатной кислоты в присутствии редкоземельных элементов)

PERIODICAL: Nauchno-tekhnicheskaya vyschey shkoly. Khimiya i khimicheskaya tekhnika, 1958, Nr 4, pp 718-720 (USSR)

ABSTRACT: A method for the rapid determination of ethylene diamine tetra acetic acid in the presence of rare earths was described. The method can be employed to separate the rare earths with ion exchange by means of  $H_4Y$ . For the determination of the  $H_4Y$  content the complexometric titration of  $H_4Y$  with calcium salt solutions was carried out with ammonium oxalate as indicator and pH 8-9. During titration the pH-value has to be regulated continuously. The initial content of  $H_4Y$  can be determined by the known quantities of the rare earth ions in the samples and by the added quantities and the back-titrated content of  $H_4Y$ . The working scheme suggested for analyses of  $H_4Y$  is applicable

Card 1/2

The Determination of Ethylene Diamino Tetra Acetic Acid in the Presence of the  
Rare Earth Elements

SOV, 1984-4-26/49

to all rare earths. There are 6 references.

ASSOCIATION: Kafedra neorganicheskoy khimii Moskovskogo gosudarstvennogo  
universiteta im. M. V. Lomonosova (Chair of Inorganic Chemistry  
at the Moscow State University imeni M. V. Lomonosov)

SUBMITTED: July 5, 1988

Card 2/2

MARTEVSKO, K L T., Cand Chem Sci -- (disc) "Chromatographic separation of rare earth elements by means of ethylenediamine tetraacetic acid." Nov, 1954, 12 pp (Disc, order no. 1611). In order of Labor ned Pomer State Univ im I.V. Leninskogo. Chem faculty.  
120 copies (kL, copy, 112)

- 17 -

MARTYNE NUKO, L.J.

PAGE : BOOK EXPLOITATION 327 402

5(2)

Ambrova and Sereb. Institute of Inorganic Materials  
Solvothermal Element Preparation, Analysis, and Properties (Rare Earth Elements)  
Preparation, Analysis, and Properties, Tsvet Moscow, 1959 531 p.  
Soviet experts presented.

Prof. Dr. I. P. Polubarnov, Professor D. N. Privalov, Dr. N. V. Kostylev, Dr. S. G. Kharlamov, Doctor I. P. Alimov, Doctor of Sciences, V. V. Shchegoleva, Doctor of Chemical Sciences, V. V. Verigin, Candidate of Chemical Sciences, N. S. Sosulin, Candidate of Chemical Sciences, N. S. Sosulin, Candidate of Chemical Sciences, and Yu. N. Salyamova, Candidate of Chemical Sciences.

This book is intended for chemists in general and for geochemists and mineralogists in particular.

Content. This collection of articles or reports prepared at the Institute of Chemical Elements suggests both the results of theoretical research and analytical chemistry. The book may be divided into three sections: methods of synthesis; the characteristics, uses, and production of new elements (rare earth elements); the methods of analyzing REE and the application of these elements. New rare elements and their structures in the glass and crystalline states, their use as catalysts, and their properties are described. The application of ion-exchange chromatography to the production of pure forms of all new earth elements. The synthesis of new REE by Dr. I. P. Polubarnov, Dr. N. A. Shchegoleva, and Dr. N. N. Salyamova. Their methods of separation are described by I. N. Bocharova. It is said to be the first comprehensive collection of methods of preparation of rare earth elements. A. V. Vinogradov, Dr. V. V. Kostylev, and G. P. Alimova, Doctor of Chemical Sciences, and V. V. Verigin, Candidate of Chemical Sciences, are concerned with the synthesis of new elements. The methods of synthesis of REE are described by Yu. N. Salyamova and P. I. Polubarnov. Their methods of separation are described by I. P. Alimov and P. I. Polubarnov. The discussion of the properties of pure products and synthesis methods is also concerned by Dr. I. P. Polubarnov. All articles are accompanied by A. N. Sogolov and his co-workers. All articles are accompanied by photographs, diagrams, tables, and bibliographic references.

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123	Polubarnov, Dr. I. P., V. S. Belogorev, V. S. Belogorev. Process of the Separation of Elements of the Cerium Sub-Group by ION Exchange
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129	Polubarnov, Dr. I. P., and N. N. Salyamova. Separation of REE by Anion Exchange
130	Soboleva, N. I., I. A. Kostylev, and V. A. Verigin. Comparative Evaluation of Electrolytic Methods of Preparing Cerium Oxide
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152	Sogolov, M. I., Dr. I. P. Polubarnov. A Rapid Method of Determining Cerium in Separates
176	Verigin, Dr. V. On the Problem of the Preparation of Cerium Oxide and the Properties of Rare Earth Elements of the Cerium Sub-Group
179	Shchegoleva, N. N., and Dr. I. P. Polubarnov. The Synthesis of a Glassy Cerium Borosilicate
186	Polubarnov, Dr. I. P. On the Separation of Cerium from the Cerium-Niobium Acid
190	Verigin, Dr. V., and Dr. I. P. Polubarnov. The Synthesis of Cerium Oxide by the Hydrolysis of Cerium Chloride
192	Polubarnov, Dr. I. P., R. S. Israels, and R. Ya. Davidzon. The Application of Ion-Exchange Chromatography on Paper for an Appropriate Separation of the Elements of the Cerium Sub-Group
199	Verigin, Dr. V. Separation of the Elements of the Cerium Sub-Group by Ion-Exchange Chromatography on Paper

5(2)

AUTHOR:

Martynenko, L. I

2010-12-14 14:54

TITLE:

The Formation of Complex Compounds of Neodymium and Ytterbium With Ethylenediamine Tetraacetic Acid on Cation-exchange Chromatography (Obrazovaniye kompleksnykh soedinenii neodima i ytterbiya s etilendiaminetetraakusnoy kislotoy pri kationoobremnoj khromatografii)

PUBLICAL:

Nauchnyye doklady vysshyey shkoly. Khimiya i khimicheskaya tekhnologiya, 1959, Nr 1, pp 84 - 86 (U.S.S.R.)

ABSTRACT:

The washing out of neodymium and ytterbium ions by means of ethylenediamine tetraacetic acid (abbreviated H<sub>4</sub>A) is observed on a cation exchanger on the basis of syntetic resin. With neodymium H<sub>4</sub>A is retained in the resin. The analysis of the compound eluted from the resin showed the complex acid H<sub>4</sub>(HdA) and its salt, respectively. Since the hexahydrate of this acid shows good solubility, the retention of H<sub>4</sub>A cannot be due to the crystallization of the acid in the resin. In this case the rate of elution was too high. Crystals of the complex acid readily tend to towards

Card 1/3

The Formation of Complex Compounds of Neodymium and Ytterbium with Ethylene-Diamine Tetraacetic Acid on Cation-exchange Chromatography

V. I. - i-a- 54

supersaturation did not separate before hours had passed after the elution from the column. Neodymium is adsorbed in the resin in the form of the "autosalt"  $\text{Nd}(\text{NdA})_x$ . In the elution with water a desorption takes place, while the "autosalt" again enters reaction in an unchanged form. The process in the individual resin layers is different; thus the concentration fluctuations of  $\text{H}_4\text{A}$  depending on the time of the adsorption column can be explained. The special tendency of  $\text{Nd}(\text{NdA})_x$  towards adsorption is to be explained by a further investigation. With ytterbium no concentration fluctuations appear in the extract, the complex compound between  $\text{Yb}^{3+}$  and  $\text{H}_4\text{A}$  obviously has no special tendency towards adsorption. This difference in the behavior of the complex compounds of neodymium and ytterbium offers an opportunity of separating the two elements by ion exchange. There are 1 figure, 1 table, and 6 references, 3 of which are Soviet.

Card 2.]

The Formation of Complex Compounds of Neodymium and Ytterbium With Ethylenediamine Tetraacetic Acid on Cation-exchange Chromatography

SGV/156-19-1-26/54

ASSOCIATION: Kafedra neorganicheskoy khimii Moskovskogo gosudarstvennogo universiteta im. M. V. Lomonosova (Chair of Inorganic Chemistry of Moscow State University imeni M. V. Lomonosov)

SUBMITTED: October 6, 1958

Card 3/3

5(2)  
AUTHORS:

Martynenko, L. I., Yeremin, G. K., Kamenev, A. I.

05891  
SOV/78-4-11-44/50

TITLE:

Chromatographic Separation of Rare Earths by Means of Tributyl Phosphate

PERIODICAL:

Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 11, p 2639 (USSR)

ABSTRACT:

The elution of the cerium group from silica gel by means of tributyl phosphate is described. Figure 1 shows that a distinct separation takes place; the content of  $\text{Me}_2\text{O}_3$  in the eluted products can attain 20 g/l which lies considerably above the concentrations usual in chromatography. As the experiment was not carried out under optimum conditions, it is to be expected that further investigations will yield even better results. There are 1 figure and 2 references.

ASSOCIATION:

Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova,  
Khimicheskiy fakul'tet, Kafedra neorganicheskoy khimii (Moscow  
State University imeni M. V. Lomonosov Chemical Department,  
Chair of Inorganic Chemistry)

SUBMITTED:

May 4, 1959

Card 1/1

YEREMIN, G.K.; KAMENEV, A.I.; MARTYNENKO, L.I.

Extraction of neodymium and praseodymium by means of some alkyl  
phosphates. Zhur.neorg.khim. 6 no.6:1487-1488 Je '61.  
(MIRA 14:11)  
(Neogymium) (Praseodymium)

KAMENEV, A.I.; MARTYNENKO, L.I.; YEREMIN, G.K.

Mechanism of the elution of macroquantities of rare earth elements by means of lactic acid. Zhur. neorg. khim. 6 no.7:1726-1727 Jl '61. (MIRA 14:7)

1. Moskovskiy gosudarstvennyy universitet imeni M.V.Lomonosova, kafedra neorganicheskoy khimii.  
(Rare earths) (Lactic acid)

MARTYNNENKO, L.I.

Certain properties of complex compounds formed by rare earths with  
ethylenediaminetetraacetic acid. Zhur.neorg.khim. 6 no.12:2704-  
2712 D '61. (MIRA 14:12)

l. Moskovskiy gosudarstvennyy universitet, kafedra neorganicheskoy  
khimii.  
(Rare earth compounds) (Acetic acid)

MITROFANOVA, N.D.; MARTYNENKO, L.I.

Composition of complex compounds formed by rare earth elements  
and nitrilotriacetic acid in the process of ion-exchange  
chromatography. Zhur.neorg.khim. 7 no.5:1049-1053 My '62.  
(MIRA 15:7)

l. Moskovskiy gosudarstvennyy universitet imeni Lomonosova,  
kafedra neorganicheskoy khimii.  
(Rare earth compounds) (Hexanoic acid)

L 11106-63  
ACCESSION NR: AP3001229

FWP(q)/EWT(m)/BDS AFETC/ASD/ESD-3 RM/JD

S/0078/63/008/006/1535/1537

59

21 57

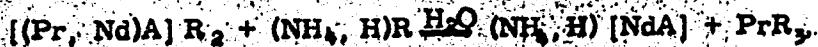
AUTHOR: Martynenko, L. I.

TITLE: Ion-exchange mechanism for separation of rare-earth elements from mixtures

SOURCE: Zhurnal neorganicheskoy khimii, v. 8, no. 6, 1963, 1535-1537

TOPIC TAGS: rare-earth element; element separation, praseodymium, neodymium, ion exchange, rare-earth group

ABSTRACT: The mechanism for separation of rare-earth elements from a mixture by means of an ion-exchange resin has been investigated, using Pr and Nd as an example. Study of the distribution of Nd and Pr between a solvent containing rare-earth ions and A<sup>-4</sup>, an acid radical of ethylenediamine tetraacetic acid [EDTA] and the resin KU-2 saturated with rare-earth ions (Ln<sup>3+</sup>) determined that adsorbed complexes acted as ion exchangers of the rare-earth elements from the solvent to the resin by the following mechanism:



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2

where R is a radical cation. The unstable complex  $[Pr, NdA]^{2+}$  decomposed on addition of an element (1% solution of  $(NH_4)_3HA$ ) to form a stable complex of Nd with EDTA, while Pr remained as a cation in the resin phase. The separation coefficient  $K_{sep}^{Pr/Nd}$  was determined from the distribution coefficients ( $K_{dist}$ ) between the resin and the solvent, and  $K_{dist}$  was determined from the ratios of the stability and instability constants ( $K_{st}$  and  $K_{inst}$ ). It was shown that in the absence of the resin, Pr and Nd are distributed between the complex form  $[LnA]^-$  and  $Ln^{+3}$ , with  $K_{sep}^{Pr/Nd} = 1.62$ , in agreement with the calculated value. In the presence of resin saturated with rare-earth ions and solvents containing  $[LnA]^-$  complex ions, the value for  $K_{sep}^{Pr/Nd}$  in transition from the static to the dynamic condition was approximately 300. It was concluded that an ion-exchange mechanism for separating rare-earth elements from a mixture can be expressed correctly only if side reactions, in particular the absorption of amino complexes  $[LnA]$  by the resin, are taken into consideration. Orig. art. has: 4 formulas and 4 tables.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova,  
Kafedra neorganicheskoy khimii (Moscow State University, Department of Inorganic  
Chemistry)

SUBMITTED: 06Dec62

DATE ACQ: 01Jul63

ENCL: 00

SUB CODE: CH,EL

NO REF Sov: 002

OTHER: 002

Card 2/2 *Py/ak*

KHARI DEV BKHARGAVA; KOVBA, L.M.; MARTYNNENKO, L.I.; SPITSYN, Vikt. I.,  
akademik

Interaction between oxides of rare-earth and alkaline earth  
metals. Dokl. AN SSSR 153 no.6:1318-1320 D '63.

(MIRA 17:1)

1. Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova.

ACCESSION NR: AP4012437

S/0078/64/009/002/0320/0329

AUTHORS: Mitrefanova, N.D.; Marty\*nenko, L.I.; Grigor'yev, A.I.

TITLE: Hydrates of the rare earth element nitrilotriacetates

SOURCE: Zhurnal neorg. khim., v. 9, no. 2, 1964, 320-329

TOPIC TAGS: rare earth nitrilotriacetate hydrate, composition, structure, rare earth nitrilotriacetate complex, high temperature hydrate, coordinate bond, ionic bond, gravimetric analysis, IR spectrum, x ray spectrum, thermogravimetric analysis, rare earth complex trihydrate, rare earth complex tetrahydrate, rare earth complex pentahydrate, hydrate bond strength, rare earth element

ABSTRACT: The rare earth element nitrilotriacetates form crystalline hydrates of different composition and structure depending on the nature of the central atom and the synthesis conditions. Complexes of the elements of the cerium subgroup and gadolinium form hydrates with 5, 4 and 3 molecules of water while the yttrium subgroup forms on the tetrahydrate. The probability of forming the

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ACCESSION NR: AP4012437

pentahydrate in the cerium subgroup decreases with the element at the end of the subgroup: samarium and gadolinium complexes form the pentahydrate only on salting out with ethanol. The water molecule bond strength is different with different types of hydrates. The aceto groups of the nitrilotriacetic acid are bonded ionically in nearly all the rare earth nitrilotriacetates. In the "high temperature" trihydrates (in which one molecule of water is especially strongly bonded), formed with the complexes of the most basic rare earth elements, La, Pr and Nd, the aceto groups form both coordinate and ionic bonds. The compounds were prepared by ion exchange, by reaction of suspensions of equivalent amounts of  $\text{Ln}_2\text{O}_3$  and nitrilotriacetic acid, by decantation, and by salting out with organic solvents. The products were analyzed gravimetrically and thermogravimetrically; X-ray and IR spectra were obtained. Orig. art. has: 10 figures and 2 tables.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova Kafedra neorganicheskoy khimii (Moscow State University, Department of Inorganic Chemistry)

Card 2/32

ACCESSION NR: AP4040518

S/0080/64/037/006/1183/1186

AUTHORS: Gorshkov, V.I.; Marty\*nenko, L.I.; Chumakov, V.A.

TITLE: Separation of rare element mixtures by continuous ionic counterflow

SOURCE: Zhurnal prikladnoy khimii, v. 37, no. 6, 1964, 1183-1186

TOPIC TAGS: rare element ionic separation, continuous ionic counterflow, rare element separation, rare element mixture separation, ionic counterflow, EDTK salt

ABSTRACT: This work was prompted by the slowness, clumsiness and low yields of the conventional method of rare element separation in batches by saturating the upper column layer of resin with a solution of rare elements in form of chlorides, while the lower, "retaining" layer of the column is saturated with copper (II), iron (III) and hydrogen ions. Through these two layers a solution of (EDTK is not explained. It may be the Russian trade name of an ion exchange resin, but it also may be the abbreviation for an organic acid) EDTK salts is circulated (most frequently tri-substituted

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ACCESSION NR: AP4040518

ammonium salt ( $\text{NH}_4$ )<sub>3</sub>HA). Instead of the above, the authors propose a continuous counterflow ion exchange method similar to that described by Spedding et al (J. Am. Chem. Soc 77, 1393 (1955)) but showing a number of advantages compared to it such as possibility of automation, simple controls, easy feed, and product separation. Two columns operating in counterflow are used. Ion exchange resin in its Cu or Cu and NH<sub>4</sub> form is fed into the upper part of column 1 where continuous frontal analysis of the mixture takes place. Yt, Gd and Sm form the more stable complexes with EDTK and therefore are accumulated in the upper part of column 1. The ion exchange resin in rare-earth-form leaving column 1 enters into column 2 against a counterflow of magnesium complexonate. Since rare element complexes with EDTK are more stable than the magnesium complexonate, the rare element ions are forced out of the resin. Pr concentrates in the lower part of column while the remainder of the solution is directed to the lower part of column 1. The ion exchanger resin in Mg-form flows from the lower part of column 2 for regeneration. This method permits visual control of separation since cation exchange resin KU-2 acquires different coloration in its various cation forms.

The method is adaptable for big scale production. The mixture to be  
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ACCESSION NR: AP4040518

separated contained 68% Nd, 20% Pr, 7% Sm, 2% Gd and other elements of the yttrium group. The equivalent height of the theoretical plate was 3.1 cm. "The authors express their gratitude to the radiological laboratory of IONKh (Inst. of Org. and Inorg. Chemistry) UkrSSR." Orig. art. has: 3 figures and 1 table

ASSOCIATION: None

SUBMITTED: 09Jul62

SUB CODE: IC

ENCL: 00

NR REF Sov: 010

OTHER: 004

Card 3/3

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CIA-RDP86-00513R001032610018-9"

MURAV'YEVA, I.A.; KOVRA, L.M.; MARTYNEVSKO, L.I.; VELIKOV, Vikt. I.

Synthesis of sodium ytterbinate. *Tr. nauch. s'ezd. po khimii*,  
1756-1757 Jl 165.

I. Kafedra neorganicheskoy khimii Moskovskogo gosudarstvennogo  
universiteta imeni M.V. Lomonosova.

L 40727-65 EPF(c)/EWT(m)/EPR/EWP(b)/EWP(t) Pr-4/Ps-4 LJP(c) JD/JG  
ACCESSION NR: AP5010581 UR/0020/65/161/003/0594/0596

AUTHOR: Bkhargava, Kh. D. i. Kovba, L. M.; Martynenko, L. I.; Spitsyn, Vikt. I. (Academician)

TITLE: New compounds of rare earths with strontium and barium oxides

SOURCE: AN SSSR, Doklady, v. 161, no. 3, 1965, 594-596

TOPIC TAGS: rare earth, strontium oxide, barium oxide, rare earth compound, strontium compound, barium compound, phase analysis, crystal structure

ABSTRACT: A series of compounds of the  $\text{SrLn}_2\text{O}_4$  or  $\text{BaLn}_2\text{O}_4$  type have been synthesized by calcining at 1100—1300°C compacted mixtures of variable ratios of corresponding oxides or salts. Reactions between strontium oxide and rare earths and between barium oxide and certain rare earths were studied by x-ray phase analysis of the reaction products. Most of the x-ray patterns of the products were obtained with RKD-57 and RKU-86 chambers. The results of phase analysis are shown in Table 1 of the Enclosure. Strontium oxide was shown to form

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L 40727-65

ACCESSION NR: AP50105B1

$\text{SrLn}_2\text{O}_4$  compounds with all rare earths from neodymium to lutetium, at a rate decreasing with an increase in the ionic radius of the rare earth element. Barium oxide reacted with the rare earths from praseodymium to lutetium, i.e., formed  $\text{BaLn}_2\text{O}_4$  compounds with the elements having a greater ionic radius than those reacting with the strontium oxide. All synthesized compounds, except  $\text{BaYb}_2\text{O}_4$ , crystallized in the rhombic system with a  $\text{CaFe}_2\text{O}_4$ -type structure. The lattice parameters of all  $(\text{Sr}, \text{Ba})\text{Ln}_2\text{O}_4$  compounds were tabulated. Apparently,  $\text{BaYb}_2\text{O}_4$  crystallized in the hexagonal system and displayed an ordered arrangement of heavy atoms, forming a superstructure. The reaction between  $\text{SrO}$  and  $\text{Sm}_2\text{O}_3$  was singled out because of a significant solubility of  $\text{SrO}$  in  $\text{Sm}_2\text{O}_3$  at 1700°C, which confirmed the possible existence of an extended homogeneity region ( $\beta$  form) in  $\text{Sm}_2\text{O}_3$ . Formation of the  $(\text{Sr}, \text{Ba})\text{Ln}_2\text{O}_4$  compounds is explained as manifestation of the amphoteric property of the rare earths. All of the compounds, except  $\text{SrYb}_2\text{O}_4$ , were found to be stable at room temperature in the presence of water vapor, moist, or dry  $\text{CO}_2$ . Orig. art. has: 4 tables.

[JK]

Card 2/4

L 40727-65

ACCESSION NR: AP5010581

ENCLOSURE: 01

Table 1. Phase analysis data

Compos- ition of the initial mixture	Cal- cina- tion temper- ature	Phase composition	Compos- ition of the initial mixture	Cal- cina- tion temper- ature		Phase composition	
				H <sub>2</sub>	O <sub>2</sub>		
SrO - La <sub>2</sub> O <sub>3</sub>	1100	13	La <sub>2</sub> O <sub>3</sub>	150	0,12	SrSm <sub>0.5</sub> O <sub>2</sub> - Sm <sub>0.5</sub> O <sub>2</sub>	
SrO - Pr <sub>2</sub> O <sub>3</sub>	1100*	12	Pr <sub>2</sub> O <sub>3</sub>	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - Pr <sub>2</sub> O <sub>3</sub>	
2SrO - Pr <sub>2</sub> O <sub>3</sub>	1100*	12	Pr <sub>2</sub> O <sub>3</sub>	250	0,12	Sm <sub>0.5</sub> O <sub>2</sub> - Pr <sub>2</sub> O <sub>3</sub>	
2SrO - Pr <sub>2</sub> O <sub>3</sub>	1100*	12	Pr <sub>2</sub> O <sub>3</sub>	250	0,12	Sm <sub>0.5</sub> O <sub>2</sub> - Pr <sub>2</sub> O <sub>3</sub>	
2SrO - Nd <sub>2</sub> O <sub>3</sub>	1100	5	SrNd <sub>0.5</sub> O <sub>2</sub> - Nd <sub>2</sub> O <sub>3</sub>	150	0,12	SrEu <sub>0.5</sub> O <sub>2</sub> - Eu <sub>2</sub> O <sub>3</sub>	
SrO - Nd <sub>2</sub> O <sub>3</sub>	1100	14	SrNd <sub>0.5</sub> O <sub>2</sub> - Nd <sub>2</sub> O <sub>3</sub>	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - Nd <sub>2</sub> O <sub>3</sub>	
SrO - Nd <sub>2</sub> O <sub>3</sub>	1100	36	SrNd <sub>0.5</sub> O <sub>2</sub> - Nd <sub>2</sub> O <sub>3</sub>	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - Nd <sub>2</sub> O <sub>3</sub>	
SrO - Nd <sub>2</sub> O <sub>3</sub>	1300	16	SrNd <sub>0.5</sub> O <sub>2</sub> - Nd <sub>2</sub> O <sub>3</sub>	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - Nd <sub>2</sub> O <sub>3</sub>	
2SrO - 15m <sub>0.5</sub> O <sub>2</sub>	1200	36	SrSm <sub>0.5</sub> O <sub>2</sub> - SrO	(e = 5,157 ± 0,001)	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - 15m <sub>0.5</sub> O <sub>2</sub>
2SrO - 25m <sub>0.5</sub> O <sub>2</sub>	1200	36	SrSm <sub>0.5</sub> O <sub>2</sub> - SrO	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - 25m <sub>0.5</sub> O <sub>2</sub>	
7SrO - 3Sm <sub>0.5</sub> O <sub>2</sub>	1200	36	SrSm <sub>0.5</sub> O <sub>2</sub> - SrO	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - 3Sm <sub>0.5</sub> O <sub>2</sub>	
8SrO - 4Sm <sub>0.5</sub> O <sub>2</sub>	1200	36	SrSm <sub>0.5</sub> O <sub>2</sub> - Sm <sub>0.5</sub> O <sub>2</sub>	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - 4Sm <sub>0.5</sub> O <sub>2</sub>	
15SrO - 6Sm <sub>0.5</sub> O <sub>2</sub>	1200	36	SrSm <sub>0.5</sub> O <sub>2</sub> - Sm <sub>0.5</sub> O <sub>2</sub>	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - 6Sm <sub>0.5</sub> O <sub>2</sub>	
15SrO - 6Sm <sub>0.5</sub> O <sub>2</sub>	1200	36	SrSm <sub>0.5</sub> O <sub>2</sub> - Sm <sub>0.5</sub> O <sub>2</sub>	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - 6Sm <sub>0.5</sub> O <sub>2</sub>	
25SrO - 8Sm <sub>0.5</sub> O <sub>2</sub>	1200	6	SrSm <sub>0.5</sub> O <sub>2</sub> - Sm <sub>0.5</sub> O <sub>2</sub>	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - 8Sm <sub>0.5</sub> O <sub>2</sub>	
15SrO - 8Sm <sub>0.5</sub> O <sub>2</sub>	1200	6	SrSm <sub>0.5</sub> O <sub>2</sub> - Sm <sub>0.5</sub> O <sub>2</sub>	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - 8Sm <sub>0.5</sub> O <sub>2</sub>	
75SrO - 8Sm <sub>0.5</sub> O <sub>2</sub>	1700	0,12	SrSm <sub>0.5</sub> O <sub>2</sub> - SrO	(e = 5,1588 ± 0,0005)	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - 8Sm <sub>0.5</sub> O <sub>2</sub>
85SrO - 8Sm <sub>0.5</sub> O <sub>2</sub>	1700	0,12	SrSm <sub>0.5</sub> O <sub>2</sub> - SrO	(e = 5,1588 ± 0,0005)	150	0,12	Sr <sub>0.5</sub> O <sub>2</sub> - 8Sm <sub>0.5</sub> O <sub>2</sub>

\* Samples were calcinated in hydrogen stream.

\*\* % of the pure SrO was found to be

5,1588 ± 0,0005 %.

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L 40727-65

ACCESSION NR: AP5010581 /

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova (Moscow State University)

SUBMITTED: 10Nov64

ENCL: 01

SUB CODE: R, GC

NO REF Sov: 003

OTHER: 002

ATD PRESS: 3231

Card 3/4

L 15610-65 EPP(n)-2/EWA(c)/EWI(m)/EWP(b)/T/EWP(t) Pr-4 IJP(c) JD/JG

UR/0020/65/161/006/1359/1361

ACCESSION NR: AP5012768

AUTHOR: Sevost'yanova, N. I.; Murav'yeva, I. A.; Kovba, L. M.; Martyenko, I. I.; Spitsyn, Vlkt. I. (Academician)

29

27

B

TITLE: New compounds of the rare earths with lithium

SOURCE: AN SSSR. Doklady, v. 161, no. 6, 1965, 1359-1361

TOPIC TAGS: rare earth compound, lithium compound, lithium terbate, lithium <sup>17</sup>  
praseodymate, lithium lanthanate, lithium ytterbate <sup>27</sup>

ABSTRACT: Two new and two known compounds of rare earths with lithium oxide have been prepared and analyzed by x-rays to supplement study of the amphoteric character of the rare earths. The new compounds were prepared by calcining at 500-800°C compacted mixtures of  $\text{Pr}_2\text{O}_3$  or  $\text{Tb}_2\text{O}_3$  and lithium carbonate. As x-ray powder patterns of the products were similar to the previously described patterns of  $\text{LiNdO}_2$  or  $\text{LiGdO}_2$ , it was possible to assign the new compounds the formula  $\text{LiPrO}_2$  or  $\text{LiTbO}_2$ . Lithium terbate ( $\text{LiTbO}_2$ ) crystallized in the rhombic system with lattice constants:  $a = 5.27 \text{ \AA}$ ,  $b = 11.16 \text{ \AA}$ , and  $c = 3.41 \text{ \AA}$ . The crystal structure of  $\text{LiPrO}_2$  is different but could not be determined from its x-ray pattern. The known compounds  $\text{LiLaO}_2$  and  $\text{LiYbO}_2$  were prepared by calcining mixtures of the nitrates of

Cord 1/2

L 15610-65

ACCESSION NR: AP5012768

2

corresponding rare earth elements with lithium nitrate.  $\text{LiLaO}_2$ , in disagreement with the earlier data, crystallized in the rhombic system and had a distorted cubic unit cell. The lattice parameters of  $\text{LiLaO}_2$  were determined. The earlier reported crystal structure of  $\text{LiYbO}_2$  was confirmed. Orig. art. has: 3 tables. [JK]

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova (Moscow State University)

SUBMITTED: 07Dec64

ENCL: 00

SUB CODE: GC, IC

NO REF Sov: 006

OTHER: 006

ATT PRESS: 1001

Card 2/2

P-1-212 MA

L 42885-66 EWT(m)/EWP(j) RM  
ACC NR: AP6020387 (A)

SOURCE CODE: UR/0192/66/007/001/0130/0131

AUTHOR: Belyayeva, K. F.; Poray-Koshits, M. A.; Mitrofanova, N. D.; Martynenko, L. I.

ORG: Moscow State University im. M. V. Lomonosov (Moskovskiy gosudarstvennyy universitet)

TITLE: X-ray structural study of neodymium nitrilotriacetate trihydrate

SOURCE: Zhurnal strukturnoy khimii, v. 7, no. 1, 1966, 130-131

TOPIC TAGS: neodymium compound, nitrogen compound, acetate, crystal structure analysis, electron density, x ray analysis

ABSTRACT: Data are presented on the lattice parameters of  $GdX \cdot 4H_2O$  and  $ErX \cdot 4H_2O$  ( $X =$  acid residue of nitriloacetic acid  $(HOOCCH_2)_3N$ ), and preliminary data on the structure of one of the two modifications of  $NdX \cdot 3H_2O$  (the so-called low-temperature modification, i. e., the trihydrate).  $GdX \cdot 4H_2O$  crystals are colorless, well-faceted hexagonal pyramids. The Laue symmetry class is  $6/mmm = D_{6h}$ , the pycnometric density  $2.31 \text{ g/cm}^3$ , and the lattice parameters  $a = 10.3$ ,  $c > 30 \text{ \AA}$ .  $ErX \cdot 4H_2O$  crystals belong to the rhombic system and are in the form of very fine rhombic prisms. The lattice parameters  $a = 12.1$ ,  $b = 21.5$ ,  $c = 9.0 \text{ \AA}$ ,  $d_{\text{calc}} = 2.40 \text{ g/cm}^3$ . Space groups  $Pna2_1$  and  $Pnam$  are possible, and  $N = 4$ . The pale-lilac, well-faceted  $NdX \cdot 3H_2O$  crystals belong to the rhombic system:  $a = 13.21$ ,  $b = 20.88$ ,  $c = 8.12 \text{ \AA}$ ,  $d_{\text{meas}} = 2.27$ ,  $d_{\text{calc}} = 2.29 \text{ g/cm}^3$ ,  $N = 8$ . Space group  $P_{bca}$ . The atomic coordinates were determined from the

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UDC: 538.736.4

L 42885-66

ACC NR: AP6020387

three-dimensional distribution of electron density, and were refined by the least-squares method. Each nitriloacetic acid residue is linked to three Nd atoms simultaneously. Orig. art. has: 1 table.

SUB CODE: 20,07/SUBM DATE: 12Aug65/ ORIG REF: 002/ OTH REF: 001

Card 2/2 bsh

GRIGOR'YEV, A.N.; MITROFANOVA, N.D.; MARTYNNENKO, L.I.

Stretching vibrations of the metal-nitrogen bond from the data  
of the infrared spectra of nitrilotriacetates. Zhur. neorg. khim.  
11 no.1:213-215 Ja '66.

(MILITARISCH)

1. Kafedra neorganicheskoy khimii Moskovskogo gosudarstvennogo  
universiteta imeni M.V.Lomonosova. Submitted March 19, 1966.

L 47185-66 EWT(m)/EWP(t)/ETI IJP(c) JD/JG  
ACC NR: AP6027193 (A) SOURCE CODE: UR/0078/66/011/008/1965/1966

AUTHOR: Ekhargava, Kh. D.; Kovba, L. M.; Martynenko, L. I.; Spitsyn, V. I.

ORG: Inorganic Chemistry Department, Moscow State University im. M. V. Lomonosov  
(Kafedra neorganicheskoy khimii, Moskovskiy gosudarstvennyy universitet)

TITLE: Reactions of barium oxide with rare earth oxides

SOURCE: Zhurnal neorganicheskoy khimii, v. 11, no. 8, 1966, 1965-1966

TOPIC TAGS: barium oxide, barium compound, rare earth compound

ABSTRACT: In a study of the solid-phase reactions of BaO with rare earth oxides, pressed pellets of stoichiometric mixtures were fired at 1000-1300°C, and the products were subjected to X-ray phase analysis with RKD-57 cameras.<sup>10</sup> The reactions of BaO with Nd<sub>2</sub>O<sub>3</sub>, Sm<sub>2</sub>O<sub>3</sub>, Gd<sub>2</sub>O<sub>3</sub> and Ho<sub>2</sub>O<sub>3</sub> proceed at a rapid rate even at 850°C, whereas the reaction with Lu<sub>2</sub>O<sub>3</sub> and especially Yb<sub>2</sub>O<sub>3</sub> takes place above 1000°C. Tm<sub>2</sub>O<sub>3</sub> and Er<sub>2</sub>O<sub>3</sub> do not react with BaO up to 1200°C; this is probably due to the fact that at this temperature the products BaTm<sub>2</sub>O<sub>4</sub> and BaEr<sub>2</sub>O<sub>4</sub> are at the verge of transition from a CaFe<sub>2</sub>O<sub>4</sub>-type structure to a BaYb<sub>2</sub>O<sub>4</sub>-type structure. BaYb<sub>2</sub>O<sub>4</sub> and BaLu<sub>2</sub>O<sub>4</sub> crystallize in the hexagonal system, and their sublattice parameters are given. The type of their superstructure could not be determined. Orig. art. has 2 tables.

SUB CODE: C7/ SUBM DATE: 15Apr65/ ORIG REF: 001/ OTH REF: 001

Card 1/1 egs

UDC: 546.65431'21

28  
B

GOLUBOV, V.I.; BAKHYNENKO, I.I.; CHIRMAKOV, V.A.

Separation of mixtures of rare-earth elements by a continuous  
countercurrent ion-exchange method. Zhur. prikl. khim. 50 no. 1  
(Zhurn. prikl. khim.)  
1976-1186 Je 102.

GRIGORYEV, V. V.

Study of some organotitanium oxides by infrared spectroscopy.  
Ph.D. neorg. khim. 1970, 10, 1212-1218.

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova,  
kafedra neorganicheskoy khimii.

L 61078-65 EPT(c)/ENG(j)/EPA(s)-2/EWT(m)/EWP(b)/EWP(t) IJP(e) JD/JG

ACCESSION NR: AP5018256

UR/0078/65/010/007/1756/1757  
546.33'668

20

19

13

AUTHOR: Murav'yeva, I. A.; Kovba, L. M.; Martynenko, L. I.; Spitsyn, V. I.TITLE: Synthesis of sodium ytterbateSOURCE: Zhurnal neorganicheskoy khimii, v. 10, no. 7, 1965, 1756-1757TOPIC TAGS: sodium ytterbate, ytterbium oxide, sodium oxide

ABSTRACT: The aim of the study was to investigate the reaction between ytterbium oxide and sodium oxide. Mixtures of the oxides in the proportions Y:Na = 1:2 and 1:3 were treated with nitric acid, the nitrate solutions were dried, and the residue was carefully heated in air until all the nitrogen oxides were driven off, then heated in a furnace for 6 hr. at 850°C. When the calcined samples were studied by x-ray phase analysis, the powder patterns revealed new lines which indicated that a reaction had taken place. It was found that the compound belongs to a hexagonal system, is isostructural with sodium ferrite and sodium indate, and hence, that its composition is  $\text{NaYbO}_2$ . The lattice constants for

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I-51078-65

ACCESSION NR: AP5018256

$\text{NaYbO}_3$  are  $a = 3.350 \pm 0.01$  Å and  $c = 16.53 \pm 0.01$  Å. Under the same conditions, no reaction was observed between ytterbium oxide and potassium oxide or rubidium oxide. Compounds were also formed by heating yttrium oxide and lanthanum oxide with sodium oxide. Orig. art. has: 1 table.

ASSOCIATION: Kafedra neorganicheskoy khimii, Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova (Department of Inorganic Chemistry, Moscow State University)

SUBMITTED: 07Dec64

RECL: 00

SUB CODE: IC

NO REF Sov: 001

OTHER: 002

Card

2/2

"APPROVED FOR RELEASE: 06/14/2000

CIA-RDP86-00513R001032610018-9

APPROVED FOR RELEASE: 06/14/2000 CIA-RDP86-00513R001032610018-9"

SEVAST'YANOVA, N.I., POGODINA, T.A., KURAYEV, M.K.; KOSTENKO, V.V.  
GLITZYN, V.YU., VYASOVSKIY, V.P.

New compounds formed by aromatic carboxylic acids with 1,3-dihydro-1,5-dioxo-2H-pyrazoles. Part 1. 1,3-dihydro-1,5-dioxo-2H-pyrazole-2-carboxylic acid and its derivatives. I. Moscow State University, Institute of Chemistry, Faculty of Chemistry, Moscow, USSR. Dokl. AN SSSR 1954(1954) No. 95, p. 185-188.

I. Moskovskiy gosudarstvennyj universitet.

KOVALEVA, A.F.; KOLOMEN, S.A.; KOCETIOVSKAYA, T.N.; LARIONOV, M.P.;  
KARTYLENKO, L.M.; SAVEL'YEV, Ye.Z.; KUZLOV, G.A., otv.  
red.; SOSKIN, A.M., red.

[Album of visual aids on economics; the section "Sosialism."  
Al'bom nagliadnykh posobii po politicheskoi ekonomii; razdel  
"Sotsializm." Leningrad, Gospolitizdat, 1960. 40 plates  
(MIRA 15:11)  
(Economics—Audio-visual aids)

L 28456-66 EMP(e)/EMT(m)/EMP(j)/T IJP(e) WW/RM/WH

ACC NR: AP6018060 (A) SOURCE CODE: UR/0020/66/168/003/0599/0602

AUTHOR: Rabinovich, I. B.; Lebedev, B. V.; Sladkov, A. M.; Kudryavtsev,  
Yu. P.; Martynenko, L. Ya.; Korshak, V. V. (Corresponding member AN SSSR)

ORG: Gorkiy State University im. N. I. Lobachevskiy (Gor'kovskiy  
gosudarstvennyy universitet); Institute of Heteroorganic Compounds,  
Academy of Sciences SSSR (Institut elementoorganicheskikh soyedineniy  
Akademii nauk SSSR)

TITLE: Carbon polymer<sup>1</sup> with increased heat capacity

SOURCE: AN SSSR. Doklady, v. 168, no. 3, 1966, 599-602

TOPIC TAGS: linear polymer, carbon polymer, chain polymer, polymer  
cross linking, carbyne, semiconducting polymer, heat capacity

ABSTRACT: The heat capacity of synthesized carbyne has been measured  
in the 80—300K range to determine the structure of this carbon polymer  
in view of the increasing interest in semiconductor<sup>15</sup> and thermal prop-  
erties of the simplest linear chain polymer with conjugated bonds<sup>7</sup> of the  
carbon polymer. Carbyne in the form of a black, fine-grain product,  
stable in air and containing 99.5% C, was synthesized by oxidation-  
polydehydrocondensation of acetylene<sup>1</sup> in the presence of bivalent copper.  
Heat capacity  $C_p$  measurements were carried out in helium atmosphere

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UDC: 541.12

L 28456-66

ACC NR: AP6018060

with 0.001° accuracy. The  $C_p$  value was accurate to 0.5%. For the purpose of comparison,  $C_p$  was also measured in Acheson graphite, C-3 domestic graphite, and acetylene black. Heat capacity was found to vary in the sequence: diamond<sup>15</sup> < graphite<sup>15</sup> < acetylene black < carbyne. Heat capacity of all nine carbyne samples was significantly higher than that of graphite, although different in each sample. This difference in  $C_p$  from one carbyne sample to another was correlated with the different ratio of the chain to lamellar structure, i.e., with partial-cross-linking of carbon chains. The samples with highest  $C_p$  were assumed to have a low degree of cross-linking, therefore to be nearly linear carbon polymers, since the value of n in the formula  $C_p = AT^B$  was nearly 1 for these samples. The n value for other samples was 1.2—1.5. Therefore, it was concluded that the products synthesized as described were different from graphite and had a lamellar-chain structure. Orig. art. has: 2 figures and 2 tables. [JK]

SUB CODE: 07/ SUBM DATE: 28Oct65/ ORIG REF: 012/ OTH REF: 007  
ATD PRESS: 5005

Card 2/2